Amendments to the Claims:

The listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

- 1. (previously presented) A recombinant, purified, or isolated polypeptide comprising an amino acid sequence selected from
- (a) the sequence of SEQ ID NO: 1;
- (b) amino acids 20 to 235 of SEQ ID NO: 1;
- (c) a sequence which has greater than 95% amino acid sequence identity with SEQ ID NO: 1; or
- (d) a sequence which has greater than 95% amino acid sequence identity with the sequence of amino acids 20 to 235 of SEQ ID NO: 1.
- 2. (previously presented) The polypeptide as claimed in claim 1 wherein the amino acid sequence of the polypeptide has greater than 97% amino acid sequence identity with SEQ ID NO:1.
- 3. (previously presented) The polypeptide as claimed in claim 1 wherein the amino acid sequence of the polypeptide has greater than 99% amino acid sequence identity with SEQ ID NO: 1.
- 4. (previously presented) The polypeptide as claimed in claim 1 wherein the amino acid sequence of the polypeptide has greater than 99% amino acid sequence identity with the sequence of amino acids 20 to 235 of SEQ ID NO: 1.
- 5. (previously presented) The polypeptide as claimed in claim 1 wherein the polypeptide consists of an amino acid sequence selected from
- (a) the sequence of SEQ ID NO: 1;

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- (b) amino acids 20 to 235 of SEQ ID NO: 1;
- (c) a sequence which has greater than 95% amino acid sequence identity with SEQ ID NO: 1; or
- (d) a sequence which has greater than 95% amino acid sequence identity with the sequence of amino acids 20 to 235 of SEQ ID NO: 1.
- 6. (previously presented) The polypeptide as claimed in claim 1 obtained from a bacterium.
- 7. (previously presented) The polypeptide as claimed in claim 1 obtained from *Mycobacterium avium* subspecies *paratuberculosis*.
- 8. (previously presented) The polypeptide as claimed in claim 1 obtained from a heterologous host transformed with a polynucleotide which encodes the polypeptide, wherein said host is capable of expressing said polypeptide.
- 9. (previously presented) The polypeptide as claimed in claim 8 wherein the host is E coli.
- 10. (previously presented) A genetic construct comprising
- (a) a promoter sequence;
- (b) an open reading frame polynucleotide encoding a polypeptide as claimed in claim 1; and
- (c) a termination sequence.
- 11. (previously presented) A recombinant, purified, or isolated polynucleotide comprising the sequence of SEQ ID NO: 2 or a variant thereof encoding a polypeptide comprising an amino acid sequence selected from
 - (a) the sequence of SEQ ID No:1;
 - (b) amino acids 20 to 235 of SEQ ID NO:1;
 - (c) a sequence which has greater than 95% amino acid sequence identity with SEQ ID NO:1; or

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- (d) a sequence which has greater than 95% amino acid sequence identity with the sequence of amino acids 20 to 235 of SEQ ID NO:1.
- 12. (original) A recombinant, purified or isolated polynucleotide with a nucleotide sequence complementary to the polynucleotide of claim 11.
- 13. (currently amended) One or more oligonucleotide or polynucleotide primers capable of <u>binding to and amplifying a polynucleotide</u> which encodes a polypeptide <u>consisting of an amino acid sequence selected from (a) the sequence of SEQ ID NO:1 or (b) amino acids 20 to 235 of SEQ ID NO:1 as claimed in claim 5 in a Polymerase Chain Reaction or other polynucleotide amplification method.</u>
- 14. (currently amended) A purified or isolated antibody capable of binding a polypeptide consisting of an amino acid sequence selected from (a) the sequence of SEQ ID NO:1 or (b) amino acids 20 to 235 of SEQ ID NO:1as defined in claim 5.
- 15. (previously presented) A vaccine composition comprising a polypeptide as claimed in claim 1 and an acceptable diluent, carrier, excipient, or adjuvant, said polypeptide being present in an amount sufficient to generate a protective immune response to *Mycobacterium avium* subspecies *paratuberculosis* infection.
- 16. (previously presented) A diagnostic composition for use in detecting the presence of *Mycobacterium avium* subspecies *paratuberculosis*, wherein said composition comprises a polypeptide as claimed in claim 1 together with one or more acceptable diluents, carriers, excipients, or adjuvants.
- 17. (previously presented) A diagnostic composition for detecting the presence of *Mycobacterium avium* subspecies *paratuberculosis*, wherein said composition comprises a polynucleotide according to claim 11 together with one or more acceptable diluents, carriers, excipients, or adjuvants.
- 18. (currently amended) A diagnostic composition for detecting the presence of *Mycobacterium avium* subspecies *paratuberculosis* comprising at least one oligonucleotide or polynucleotide primer capable of amplifying a polynucleotide which encodes a polypeptide as claimed in claim 5 in a Polymerase Chain Reaction or other

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polynucleotide amplification method, wherein the polypeptide consists of an amino acid sequence selected from (a) the sequence of SEQ ID NO:1 or (b) amino acids 20 to 235 of SEQ ID NO:1.

- 19. (previously presented) A diagnostic composition for detecting the presence of *Mycobacterium avium* subspecies *paratuberculosis* comprising an antibody according to claim 14 together with one or more acceptable diluents, carriers, excipients, or adjuvants.
- 20. (currently amended) A method of detecting Johne's disease including preclinical Johne's disease in an animal comprising contacting either the animal or a sample from the animal with a polypeptide as elaimed in claim 5 and detecting an immune response indicative of the presence of *Mycobacterium avium* subspecies *paratuberculosis*, the polypeptide consists of an amino acid sequence selected from (a) the sequence of SEQ ID NO:1 or (b) amino acids 20 to 235 of SEQ ID NO:1.
- 21. (previously presented) The method according to claim 20 wherein the response is a delayed-type hypersensitivity response.
- 22. (currently amended) The method according to claim 20 wherein said detecting comprises detecting the presence of antibodies that bind the polypeptide as claimed in claim 5.
- 23. (previously presented) The method according to claim 22 wherein the detection of the presence of antibodies is by ELISA, radioimmunoassay or Western blotting.
- 24. (currently amended) A method of detecting Johne's disease including preclinical Johne's disease in an animal, the method comprising contacting a sample from the animal either with a purified or isolated antibody capable of binding a polypeptide consisting of an amino acid sequence selected from
- (a) the sequence of SEQ ID NO:1 or
- (b) amino acids 20 to 235 of SEQ ID NO:1 or

with a composition comprising an antibody specific to a polypeptide consisting of an amino acid sequence selected from

- (a) the sequence of SEQ ID NO:1 or
- (b) amino acids 20 to 235 of SEQ ID NO:1as claimed in claim 5,

and one or more acceptable diluents, carriers, excipients, or adjuvants, and detecting a polypeptide which binds to the antibody.

- 25. (previously presented) The method according to claim 24 wherein the presence of bound antibody is determined by ELISA, radioimmunoassay or Western blotting.
- 26. (previously presented) The method according to claim 24 for detecting the presence of *Mycobacterium avium* subspecies *paratuberculosis* at a preclinical phase of Johne's disease.
- 27. (currently amended) A method of detecting Johne's disease including preclinical Johne's disease in an animal comprising contacting a sample from the animal with a composition comprising at least one oligonucleotide or polynucleotide primers capable of binding to and amplifying a polynucleotide which encodes a polypeptide consisting of an amino acid sequence selected from (a) the sequence of SEQ ID NO:1 or (b) amino acids 20 to 235 of SEQ ID NO:1 as claimed in claim 4 in a polynucleotide amplification method and detecting the amplification product.
- 28. (previously presented) The method as claimed in claim 27 wherein the polynucleotide amplification method is a polymerase chain reaction method.
- 29. (previously presented) The method according to claim 27 for detecting the presence of *Mycobacterium avium* subspecies *paratuberculosis* at a preclinical phase of Johne's disease.
- 30. (currently amended) A method of detecting Johne's disease in an animal comprising contacting a sample from the animal with a composition comprising a polynucleotide capable of binding to a polynucleotide which encodes a polypeptide consisting of an amino acid sequence selected from (a) the sequence of SEQ ID NO:1 or (b) amino acids 20 to 235 of SEQ ID NO:1as claimed in claim 5.
- 31. (previously presented) The method according to claim 30 wherein said polynucleotide is detectably labeled.
- 32. (previously presented) The method according to claim 31 wherein said detectable label is a radioisotope or fluorescent tag.

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- 33. (previously presented) A method of prophylactically or therapeutically treating an animal against Johne's disease which comprises administering to an animal a polypeptide as claimed in claim 1 to produce a protective immunological response in the animal.
- 34. (previously presented) The method according to claim 33 which is a therapeutic method.
- 35. (previously presented) The method according to claim 33 which is a prophylactic method.
- 36. (original) A method of vaccinating against Johne's disease which comprises administering to an animal a vaccine composition as claimed in claim 15 in an amount sufficient to produce a protective response.
- 37. (previously presented) The method according to claim 36 wherein said administration is performed on a single occasion.
- 38. (previously presented) The method according to claim 36 wherein said administration is performed on more than one occasion.
- 39. (previously presented) The method as claimed in claim 36 wherein $0.1-1000\mu g/Kg$ is administered of a recombinant, purified, or isolated polypeptide comprising an amino acid sequence selected from
- (a) the sequence of SEQ ID NO: 1;
- (b) amino acids 20 to 235 of SEQ ID NO:1;
- (c) a sequence which has greater than 95% amino acid sequence identity with SEQ ID NO:1; or
- (d) a sequence which has greater than 95% amino acid sequence identity with the sequence of amino acids 20 to 235 of SEQ ID NO:1.
- 40. (previously presented) The method as claimed in claim 39 wherein $5-500\mu g/Kg$ of the polypeptide is administered.
- 41. (currently amended) A kit for use in detecting the presence of *Mycobacterium* avium subspecies paratuberculosis comprising at least two of the following:

a polypeptide as claimed in claim 5;

an antibody that binds a said-polypeptide consisting of an amino acid sequence selected from (a) the sequence of SEQ ID NO:1 or (b) amino acids 20 to 235 of SEQ ID NO:1; or a reagent for determining antigen-antibody binding.

- 42. (previously presented) A host cell transformed with a polynucleotide of claim 11.
- 43. (original) A vector comprising the construct as claimed in claim 10.
- 44. (original) A host cell incorporating a construct of claim 10.
- 45. (original) A host cell incorporating a vector as claimed in claim 43.
- 46. (previously presented) The host cell according to claim 45 wherein said vector exists within the host cell as a plasmid.
- 47. (previously presented) The host cell according to claim 45 wherein said vector is integrated into the genome of the host cell.
- 48. (previously presented) The method as claimed in claim 20 wherein the animal is a ruminant.
- 49. (previously presented) The method as claimed in claim 48 wherein the animal is a sheep.
- 50. (previously presented) The method as claimed in claim 33 wherein the animal is a ruminant.
- 51. (previously presented) The method as claimed in claim 50 wherein the ruminant is a sheep.
- 52. (cancelled)